



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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| (22) International Filing Date: 4 August 1997 (04.08.97)   |  |  |   |  |  |
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| (72) Inventor; and   |  | <b>Published</b>   |   |  |  |
| (75) Inventor/Applicant (for US only): GUANDALINI, Stefano [IT/IT]; Via Napoli, 253, I-80018 Mugnano Di Napoli (IT).   |  | Without international search report and to be republished upon receipt of that report.   |   |  |  |
| (74) Agent: SARPI, Maurizio; Studio Ferrario, Via Collina, 36, I-00187 Roma (IT).  |  |  |   |  |  |
| <b>(54) Title:</b> TREATMENT OF THE ACUTE INFANT'S DIARRHOEA AND PREVENTION OF ALLERGIC REACTIONS TO FOODS SWALLOWED IN THE FOLLOWING PHASE BY ADMINISTERING LACTOBACILLUS GG IN THE ORAL REHYDRATING SOLUTION   |  |  |   |  |  |
| <b>(57) Abstract</b>   |  |  |   |  |  |
| <p>The early administration of Lactobacillus GG during the rehydrating phase is capable of shortening the duration of the diarrhoea, preventing the following food allergy syndrome in the patient, and promoting a faster weight increase. To this purpose there is provided a preparation to be administered by mouth and formed of an oral rehydrating solution (ORS) of the commercial type, such as Dicodral 60, in which an effective amount of both alive and inactivated ferments Lactobacillus GG is contained.</p> |  |  |   |  |  |

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Treatment of the acute infant's diarrhoea and prevention of allergic reactions to foods swallowed in the following phase by administering Lactobacillus GG in the oral rehydrating solution

The present invention relates to the treatment of acute infant's diarrhoea and more particularly the early use of the bacterium Lactobacillus GG during the rehydrating phase with the purpose of reducing the duration of the 05 diarrhoea, preventing food allergy syndromes in the patient, and promoting a faster weight recovery.

Acute diarrhoea due in most cases to intestinal infections (gastroenteritis) acquired by orofecal way is still a very 10 important sanitary problem. In fact, 2 to 3 billions of cases of acute diarrhoea occur yearly in the world, such cases causing an estimated mortality of about 5 million infants aged up to five.

In Italy and West Europe the mortality due to 15 gastroenteritis has gone progressively and considerably down in recent years and is reduced to 1-10 cases over 100,000 infants/year today. However, also under such circumstances, acute diarrhoea is an event of great importance considering that it is very frequent (recent 20 epidemiological studies in Scandinavia and in Italy show that each infant has, as an average, a little more than one episode/year of acute diarrhoea). Therefore, direct cost for the treatment and indirect cost due to the absence of the mothers from work are huge. It should not 25 be forgotten that in 4-6% of the cases an acute diarrhoea lasts more than 14 days, thus exposing the infant to the

real risk of developing malnutrition or acquiring food allergies (which actually take place in a large number of cases), thus inducing further complications and, in the best case, excluding some foods for very long time.

05 As can be seen from the foregoing, it should be very useful to have therapeutical means capable of reducing the duration of the diarrhoeic symptomatology, blocking effectively the possible evolution regarding the food allergy, and improving at the same time the digestion-  
10 absorption conditions in order to promote the weight recovery of the little patients.

The present invention seeks to provide a formulation responsive to such requirements.

15 The current therapy of gastroenteritis is addressed to an unquestionable physiopathologic approach: administering an oral rehydrating solution (ORS) based on a mixture of electrolytes and glucose formulated according to precise  
20 requirements so as to promote the intestinal absorption of water and mineral salts and to restore the hydro-electrolytic wealth endangered by diarrhoea. Such an approach, which has been made possible by the knowledge of the physiopathology of the intestinal absorption-digestion  
25 processes and has been widely used for the last 10-15 years also in the adult's diarrhoea, has certainly contributed to save many human lives and cannot be questioned at all. However, as well known, it takes no effect on the symptoms, the diarrhoea persisting  
30 unchanged, and then is only partially responsive to the above-mentioned requirements.

It is also known that apart from some specific substances the etiologic therapy of the gastroenteritis cannot be carried out as only some bacterial enteric pathogens may be eradicated by a specific therapy accompanied by a  
05 shortening of the symptomatology.

In recent years it has been proved (Raza et al., 1995) that a milk enzyme, Lactobacillus casei subspecies rhamnosius (so-called also Lactobacillus GG initially isolated from the intestinal bacterial flora of a man),  
10 administered both in the form of powder and yoghurt produced by the fermentation of the milk with said enzyme, is capable of shortening by about 2 days the duration of the diarrhoea in little patients with gastroenteritis by rotavirus which is considered as the most common etiologic  
15 agent of acute infant's diarrhoea in the world. It has been also proved that the administering of such an enzyme has no side-effects at all and is accompanied by an increase of the immunity to rotavirus.

Furthermore, it has been thereafter proved that the same  
20 product is also capable of thoroughly preventing the increase of the intestinal permeability of the rats caused by the early administering of milk proteins of different species to the growing animals. The latter result is of great importance: it is well known that during an  
25 infectious gastroenteritis (that by rotavirus is really the most studied condition) the permeability of the intestinal mucosa is altered by the damage induced by the intestinal infectious process. That such a condition predisposes to the entrance of heterologous food  
30 macromolecules and then is capable to initiate a food allergy syndrome in susceptible subjects is an

incontrovertible proof. In fact, it is well supported with documentary evidence that a considerable amount of the allergies to cow's milk afflicting about 3% of the infants came into being after a gastroenteritis.

05

Starting from such remarks it is assumed that the early administering of Lactobacillus GG during the rehydration phase (i.e. the initial acute diarrhoea treatment phase in which the oral rehydrating solution (ORS) is administered 10 without other foods) can not only further reduce the duration of the diarrhoea (allowing a better and earlier contact of the agent with the damaged intestinal mucosa) but also allow the allergy to food (especially proteins of cow's milk) administered in the following phase to be 15 prevented.

The result of an experimentation carried out on a sample of several little patients proved that a preparation to be administered by mouth and formed of the ferment Lactobacillus GG in a rehydrating solution, identified as 20 the commercial preparation Dicodral 60, in an amount of 100,000 to 1,000,000,000 C.F.U. (Colony Forming Unit) every 500 ml solution, prevents food allergy syndromes besides strongly influencing the duration of the diarrhoea, and allows the weight to be faster recovered.

25

The effectiveness of administering the above-mentioned preparation is self-evident from the results of the experimentation shown in Tables 1 and 2.

Children aged one to twelve having 4 evacuations/day of 30 liquid faeces for one to some days but not more than 5 days were subjected to test. Criteria for the exclusion

from the test were: preceding treatment with antidiarrhoea products, syndrome of short intestine, associated renal or hepatic diseases, paralytic ileum, chronic inflammatory intestine diseases (Crohn disease, ulcerous rectocolitis).

05 The test was conducted according to a double-blind method controlled by placebo. The enlisted patients were randomly assigned either to group A assuming Lactobacillus GG (added to the oral rehydrating solution (ORS) Dicodral 60: 10,000,000 C.F.U./250 ml ORS) or to group B assuming 10 placebo (only Dicodral 60). All patients were rehydrated with said solutions for 6-8 hours and then resumed the usual diet still assuming the oral rehydrating solution until the end of diarrhoea. Furthermore, registry, anthropometric, clinic data of each patient was registered 15 as well as a sample of faeces was taken for the analysis of the following enteric pathogens: Rotavirus and Adenovirus of the enteric type (40-41), Giardia and Cryptosporidium, Salmonella, Shigella, Campylobacter, Yersinia, Aeromonas, E. Coli pathogens (ETEC, EPEC, EHEC, 20 EIEC, EAEC, searched by specific DNA probes). Finally, analysis were carried out on the following parameters which were registered at the beginning and at periodic intervals during the analysis: body weight, defecation (frequency and quality), total duration of diarrhoea, oral 25 feeding and complications such as vomit and intolerance to carbohydrates (the latter searched daily by Clinitest).

#### RESULTS

Data relates to 32 patients, 17 in group A, 15 in group B. Both groups were comparable in age, weight, and duration 30 of diarrhoea at the admission (table 1). The medium age was 23.9 months (range: 2-61) in group A, 25.2 months

(range: 3-96) in group B. At the admission diarrhoea was lasting on the average from 2.9 days (range: 1-5) in group A, 2.6 days (range: 1-4) in group B. The mean duration of diarrhoea from the beginning of the treatment was 1.3 days  
05 (range 1-3) in group A, and 1.4 days (range 1-3) in group B. The mean duration of diarrhoea from the beginning of the treatment could be estimated in hours for 15 patients: it was 17.1 hours (range: 4-34) in group A, and 27 hours (range: 6-70) in group B (table 2). The mean number of  
10 liquid evacuations in the two days of evaluation was 5.6 faecal discharges (range 1-11) in group A and 7.3 faecal discharges (range 1-24) in group B. The mean weight increase was always greater in group A in all of the evaluation days as shown in the diagram. Among the  
15 complications, vomit was present in 17.6% of group A and 20% of group B; the intolerance of lactose was present in 5.8% of group A and 6.6% of group B. Finally, among the isolated pathogens, Rotavirus was isolated in 11.76% of group A and 6.6% of group B.  
20 The results of the experiments have to be considered as preliminary, however, they allow as from now some salient points to be pointed out:  
Lactobacillus GG administered by mouth in the oral rehydrating solution has proved to be well tolerated and  
25 did not result in any intolerance or side effects.  
The duration of an acute diarrhoea tends to be shorter in the treated patients than in the controls, especially if the duration is expressed in hours; however, the dispersion of the values and the little number of cases do  
30 not allow yet to confirm such a statement according to a statistic validation.

The most interesting data, however, is the faster weight increase of the infants assuming the ferment than the controls. As the latter data relates to the first 24 hours of observation, it is of course sign of a more effective 05 rehydration and/or re-nourishing. One could then suppose that the treated patients markedly improve their digestion conditions after a few hours administration with the consequence of a faster restoration of the hydro-electrolytic wealth.

10 Table 1.- Observation Data

|                                     | Group A      | Group B        | P    |
|-------------------------------------|--------------|----------------|------|
|                                     | 1-GG (n=17)  | Placebo (n=15) | N.S. |
| Age (month)                         | 23.9 (2-61)  | 25.2 (3-96)    | N.S. |
| Weight (g)                          | 11841 ± 6250 | 13439 ± 6700   | N.S. |
| 15 Undernourished                   | 2 (12%)      | 2 (13%)        | N.S. |
| No dehydration                      | 7%           | 7%             | N.S. |
| Dehydration < 5%                    | 93%          | 93%            | N.S. |
| Dehydration 5-10%                   | -            | 14%            | N.S. |
| Dehydration > 10%                   |              |                | N.S. |
| 20 Duration of the diarrhoea (days) | 2.9 ± 0.7    | 2.6 ± 1.1      | N.S. |
| Rotavirus diarrhoea                 | 2 (11.76%)   | 1 (6.6%)       | N.S. |

Table 2.- Clinic Course

|                        | Group A (n=17) | Group B (n=15) |
|------------------------|----------------|----------------|
| Vomit (%)              | 3 (17.64%)     | 3 (20%)        |
| Intolerance to lactose | 1 (5.88%)      | 1 (6.64%)      |
| Duration of diarrhoea  |                |                |
| from the beginning of  | 1.33 days      | 1.46 days      |
| 30 the treatment       | range (1-3)    | range (1-3)    |

|  | Group A (n=9) | Group B (n=6) |
|--|---------------|---------------|
|--|---------------|---------------|

Duration of diarrhoea

|  |                            |                          |
|--|----------------------------|--------------------------|
| from the beginning of<br>the treatment | 17.1 hours<br>range (4-34) | 27 hours<br>range (6-70) |
|--|----------------------------|--------------------------|

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As can be seen, the product of the present invention may then find application in the acute infant's diarrhoea, in the infectious infant's gastroenteritis, in the therapy and prevention of the infant's protracted diarrhoea 10 syndrome as well as in the prevention of the allergy to cow's milk caused by gastroenteritis.

## Claims

1. Use of an effective amount of both alive and inactivated ferments Lactobacillus GG in the oral rehydrating solution (ORS) for the early treatment of the acute infant's diarrhoea of a variety of aetiologies.

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2. A preparation to be administered by mouth for the treatment of the acute infant's diarrhoea which is formed of a rehydrating solution (ORS) containing an effective amount of both alive and inactivated Lactobacillus GG.

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3. A preparation to be administered by mouth for the treatment of the acute infant's diarrhoea which is formed of a rehydrating solution (ORS) containing an amount of both alive and inactivated Lactobacillus GG between 15 100,000 and 1,000,000,000 C.F.U. (Colony Forming Units) every 500 ml solution.

4. The preparation to be administered by mouth for the treatment of the acute infant's diarrhoea of the preceding 20 claim, wherein the rehydrating solution (ORS) is identified by the commercial preparation Dicodral 60.

5. A preparation to be administered by mouth for the treatment of the acute infant's diarrhoea of claim 4, 25 wherein the amount of Lactobacillus GG is 10,000,000 C.F.U. every 250 ml Dicodral 60.

6. A preparation to be administered by mouth for the treatment of the acute infant's diarrhoea of claims from 2

on, wherein it reduces the duration of the diarrhoea, prevents allergies to foods swallowed in the following phase, and allows a faster weight increase.

05 7. A method for the treatment of acute infant's diarrhoea, the prevention of allergies to foods swallowed in the following phase and a faster weight increase, wherein there is provided the early administration of Lactobacillus GG in the initial phase of treatment of the  
10 acute diarrhoea by adding it to the oral rehydrating solution (ORS).



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| (51) International Patent Classification <sup>6</sup> :<br><br>A61K 35/74  |  | A3   | (11) International Publication Number: <b>WO 98/06411</b><br><br>(43) International Publication Date: 19 February 1998 (19.02.98) |
| (21) International Application Number: PCT/IT97/00201<br><br>(22) International Filing Date: 4 August 1997 (04.08.97)<br><br>(30) Priority Data:<br>RM96A000571 9 August 1996 (09.08.96) IT  |  | (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). |   |
| (71) Applicant (for all designated States except US): DICOFARM S.P.A. [IT/IT]; Via F.S. Nitti, 11, I-00191 Roma (IT).<br><br>(72) Inventor; and<br><br>(75) Inventor/Applicant (for US only): GUANDALINI, Stefano [IT/IT]; Via Napoli, 253, I-80018 Mugnano Di Napoli (IT).<br><br>(74) Agent: SARPI, Maurizio; Studio Ferrario, Via Collina, 36, I-00187 Roma (IT). |  | <b>Published</b><br><i>With international search report.</i><br><i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>  |   |
|  |  | (88) Date of publication of the international search report: 7 May 1998 (07.05.98)   |   |

(54) Title: USE OF LACTOBACILLUS GG IN THE REHYDRATING SOLUTION

## (57) Abstract

The early administration of Lactobacillus GG during the rehydrating phase is capable of shortening the duration of the diarrhoea, preventing the following food allergy syndrome in the patient, and promoting a faster weight increase. To this purpose there is provided a preparation to be administered by mouth and formed of an oral rehydrating solution (ORS) of the commercial type, such as Dicodral 60, in which an effective amount of both alive and inactivated ferments Lactobacillus GG is contained.

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# INTERNATIONAL SEARCH REPORT

Intell. 1st Application No

PCT/IT 97/00201

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 A61K35/74

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

| Category | Citation of document, with indication, where appropriate, of the relevant passages  | Relevant to claim No. |
|----------|---|-----------------------|
| X, P     | <p>MINNA KAILA ET AL: "NUTRITIONAL MANAGEMENT OF ACUTE DIARRHEA"<br/>           NUTRITION TODAY,<br/>           vol. 31, no. 6, November 1996 - December 1996,<br/>           pages 16S-18S, XP002058044<br/>           see the whole document</p> <p>---</p> <p>WO 91 15199 A (MEDICIS CORP) 17 October 1991<br/>           see page 10, line 5 - page 13, line 18</p> <p>---</p> <p>-/-</p> | 1-7                   |
| X        |   | 1                     |

Further documents are listed in the continuation of box C.

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| 6 March 1998   | 25.03.1998   |
| Name and mailing address of the ISA<br>European Patent Office, P.B. 5818 Patenttaan 2<br>NL - 2280 HV Rijswijk<br>Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,<br>Fax: (+31-70) 340-3016 | Authorized officer<br><br>Rempp, G                 |

## INTERNATIONAL SEARCH REPORT

International Application No  
PCT/IT 97/00201

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

| Category | Citation of document, with indication, where appropriate, of the relevant passages  | Relevant to claim No. |
|----------|---|-----------------------|
| X        | MINNA KAILA ETAL.: "VIABLE VERSUS INACTIVATED LACTOBACILLUS STRAIN GG IN ACUTE ROTAVIRUS DIARRHOEA." ARCHIVES OF DISEASE IN CHILDHOOD, vol. 72, no. 1, January 1995, pages 51-53, XP002058045<br>see the whole document<br>---                | 1-7                   |
| X        | A. R. PANT ET AL.: "LACTOBACILLUS GG AND ACUTE DIARRHOEA IN YOUNG CHILDREN IN THE TROPICS." JOURNAL OF TROPICAL PEDIATRICS, vol. 42, no. 3, June 1996, pages 162-165, XP002058046<br>see the whole document<br>---                            | 1                     |
| X        | HELI MAJAMAA ET AL.: "LACTIC ACID BACTERIA IN THE TREATMENT OF ACUTE ROTAVIRUS GASTROENTERITIS." JOURNAL OF PEDIATRIC GASTROENTEROLOGY AND NUTRITION, vol. 20, no. 3, April 1995, pages 333-338, XP002058047<br>see the whole document<br>--- | 1                     |
| A        | EP 0 199 535 A (NEW ENGLAND MEDICAL CENTER INC) 29 October 1986<br>---  |                       |
| A        | US 5 413 785 A (NANJI AMIN A) 9 May 1995<br>---   |                       |
| A        | WO 94 18997 A (VALIO LTD ;GORBACH SHERWOOD L (US)) 1 September 1994<br>---  |                       |
| A        | EP 0 271 364 A (BIOREM C C) 15 June 1988<br>-----   |                       |

# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/IT 97/00201

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1.  Claims Nos.:  
because they relate to subject matter not required to be searched by this Authority, namely:  
see FURTHER INFORMATION sheet PCT/ISA/210
2.  Claims Nos.:  
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3.  Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1.  As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2.  As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3.  As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4.  No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

### Remark on Protest

- The additional search fees were accompanied by the applicant's protest.
- No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

Claims Nos.: 7

because they relate to subject matter not required to be searched by this Authority, namely:

Rule 39.1(iv) PCT - Method for treatment of the human or animal body by therapy

Remark : Although claim 7 is directed to a method of treatment of the human/animal body , the search has been carried out and based on the alleged effects of the compound/composition.

## INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

/IT 97/00201

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